

Is there a gender difference in antidiuretic response to desmopressin in children?

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ABSTRACT

Women with nocturia are more sensitive to desmopressin, a synthetic arginine vasopressin (AVP) analogue, with significant antidiuretic responses to desmopressin orally disintegrating tablet (ODT) 25 µg, compared with men who require 58 µg to achieve similar responses. In children the current desmopressin dose recommendation to treat primary nocturnal enuresis (PNE) is the same for boys and girls. This *post hoc* analysis of data from a randomised, double-blind single-dose study of 84 children with PNE aged 6 - 12 years explored gender differences in sensitivity to desmopressin in children. Following water loading to suppress endogenous AVP, placebo or desmopressin 30, 60, 120, 240, 360 or 480 µg was administered when urinary production reached >0.13 mL/min/kg. The endpoints of urinary osmolality and duration of urinary-concentrating action (DOA) (above three thresholds: 125, 200 and 400 mOsm/kg) were analysed to compare efficacy in boys and girls, in each treatment group. The DOA and urinary osmolality were similar in both genders in the desmopressin 120 - 480 µg groups. Boys receiving desmopressin ODT 30 - 60 µg tended to increased urinary osmolality and experienced 1 - 2 hours longer DOA than girls. The same pattern of higher values in boys compared with girls was seen for all measures of urinary osmolality. **Conclusion:** In a limited sample of pre-pubertal children the antidiuretic response to desmopressin was largely similar between genders, in contrast to findings in adults.

Keywords: Desmopressin; Antidiuresis; Gender

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Difference; Paediatric; Arginine Vasopressin Receptor 2

1. INTRODUCTION

The concept of a gender difference in renal sensitivity to desmopressin has been supported by several studies in adult patients with nocturia. In the double-blind, placebo-controlled NOCTUPUS studies, testing desmopressin tablets (0.1 mg, 0.2 mg or 0.4 mg) in 144 women [1] and 151 men [2], a ≥50% reduction in nocturnal voids was achieved in 46% of women and in 34% of men; women had a 78% increase in mean first sleep period compared with a 59% increase in men. In a trial of desmopressin orally disintegrating tablets (ODT; Minirin® Melt), 799 nocturia patients were randomised to placebo or desmopressin ODT (10, 25, 50 or 100 µg), for one month [3]. *Post hoc* analyses by gender demonstrated a significantly greater decrease in nocturnal urine volume in women at the lower doses of 10 µg and 25 µg compared with men (ED₅₀ 16.1 µg vs. 43.2 µg), suggesting significantly higher sensitivity to desmopressin in women [4]. A recent pharmacodynamic (PD) study of desmopressin in water-loaded Japanese patients with nocturia found that male patients required 58 µg desmopressin ODT to achieve a similar duration of antidiuretic action as achieved by female patients receiving desmopressin 25 µg [5]. A recent meta-analysis of data from nocturia trials with desmopressin demonstrated a durable gender difference in sensitivity to desmopressin in favour of females. Greater efficacy was observed with desmopressin ODT 25 µg in females compared with males, which endured over the long term (one year or more) [6]. Two physiological mechanisms behind this gender difference in renal sensitivity to desmopressin have been suggested, including the genetic X-inactivation mechanism and the

endogenous regulation of AVP by sex hormones [4].

In children with monosymptomatic nocturnal enuresis (MNE), current desmopressin ODT dosing recommendations (60 - 240 µg) are the same for boys and girls. However, there is substantial evidence that absence of a nocturnal increase in AVP release is a contributing factor for nocturnal polyuria and enuresis [7], and that more boys than girls are affected by the condition. In a large epidemiological study, 16.1% of children aged five years were reported to have PNE, with double the proportion of boys compared with girls, 20.7% vs. 10.8%, respectively, reported to have PNE [8]. Several research papers have confirmed that females have significantly lower basal AVP levels compared to males, both in children [9] and in young adults [10-12]. Since no difference in urine volume or osmolality was found in these studies, it is reasonable to assume that women and girls concentrate urine at lower plasma AVP levels than men and boys.

Emerging evidence supports the hypothesis that females have an increased sensitivity to both AVP and to the AVP receptor 2 (V2 receptor) specific agonist, desmopressin. In adult patients with nocturia, gender-specific dosing of desmopressin provides the clinical advantage of minimising the risk of hyponatraemia while still achieving a meaningful patient benefit in reducing nocturnal voids [13]. However, there is a lack of evidence of gender differences in children in response to V2 receptor stimulation, and no paediatric gender-specific dose recommendations have been suggested.

Thus, the purpose of this paper is to explore possible gender differences in children by quantifying the urine-concentrating effects of desmopressin at different dose-levels using a gender-stratified analysis of a previously reported clinical study in children. While we recognise that a *post hoc* analysis is not as powerful as a primary analysis, ethical considerations led us to explore relevant data already collected prior to setting up a new interventional study in children. The current analysis also supports power calculations for any prospective study protocols in a paediatric population examining gender difference in response to V2 receptor stimulation.

2. PATIENTS AND METHODS

Clinical data were derived from a PD double-blind, randomised, parallel-group study in children and adolescents with primary MNE [14].

2.1. Patients

The study was conducted in 84 children and adolescents. Boys or girls aged 6 - 13 years and of normal body weight (body mass index [BMI] 14 - 25 kg/m²) were eligible for the study if primary MNE had been demon-

strated during a run-in period. All patients were otherwise healthy. Exclusion criteria included evidence of organic urological disease, diurnal urinary incontinence, diabetes insipidus, ongoing urinary tract infection or any clinically significant disease likely to interfere with the evaluation; use of antibiotics, diuretics or any drug affecting urinary concentration; use of any therapy (pharmacological or conditioning), including desmopressin for PNE within 30 days of study entry; and abnormalities or disease of the oral cavity that might affect the release or absorption of the orally dispersible formulation.

2.2. Study Design

The study design and the clinical and safety results of this study have been reported in detail elsewhere [14]. Briefly, eligible patients were randomised to either placebo or one of six single doses of 30, 60, 120, 240, 360 or 480 µg desmopressin. All patients were water-loaded before dosing to suppress endogenous AVP. Desmopressin or placebo were administered when urinary production reached >0.13 mL/min/kg. Urinary volume, osmolality and duration of urinary-concentrating action (DOA) (above three threshold levels: 125, 200 and 400 mOsm/kg) were measured.

The study was approved by the institutional review board or ethics committee for each site, the Declaration of Helsinki was followed, and informed consent obtained from all patients and healthy volunteers. All data used are based on the ODT formulation of desmopressin.

2.2.1. Software

Statistical evaluation was performed using SAS (v9.2).

2.2.2. Endpoints and Analysis

The mean, maximum and area under curve (AUC) of urinary osmolality and the DOA (above three threshold levels: 125, 200 and 400 mOsm/kg) were compared between genders, for each treatment group, unless $n \leq 1$. For DOA analysis, patients with urine osmolality above threshold level according to these definitions, at the end of the measurement period, were censored at the time their overhydration procedure stopped. Plots of the distributions of time-adjusted AUC of osmolality, the AUC of urinary osmolality for ten hours following desmopressin administration and maximum urinary osmolality were produced.

Due to the limited sample size and skewed gender distribution of PNE (the ratio of boys to girls was 3.2:1), no formal statistical testing was done. Any comparison between the two genders is therefore based on descriptive statistics, supplemented by visual evaluation of the graphs.

3. RESULTS

3.1. Demographics

There were no major demographic differences in the baseline characteristics of patients in each of the treatment groups (**Table 1**). Since no prospective gender stratification was planned, and PNE is more prevalent in boys, the study had a skewed gender distribution with more boys than girls in all treatment groups. The majority of patients were of a pre-pubertal age, with a mean age of nine years (range 6 - 13). No patient was overweight according to the definition of BMI above 25 kg/m².

3.2. Duration of Urinary-Concentrating Action above Three Threshold Levels: 125, 200 and 400 mOsm/kg

In the higher dose groups, no clear difference between the genders in DOA was seen, although direct compari-

son between genders in the 120 and 240 µg dose groups was prevented as only one girl was assigned to each of those dose groups. Only in the highest dose group (desmopressin 480 µg) was a modestly longer mean DOA observed in girls compared with boys. In the low-dose groups (desmopressin 30 and 60 µg) boys had 1.6 - 2 hours longer mean DOA compared with girls for all three threshold levels of osmolality (**Table 2**).

Examining the two lowest dose groups together by gender produced a DOA distribution curve for boys that was rightward shifted compared with the DOA distribution curve for girls, indicating a modest increase in mean DOA of desmopressin in boys, due in part to more outliers amongst the boys with a DOA of around 10 - 12 hours (**Figure 1**).

3.3. Mean, Maximum and AUC of Urinary Osmolality

In the higher dose groups all three measurements of uri-

Table 1. Demographic characteristics.

Dose (µg)	n		Age (years)		BMI (kg/m ²)	
	Female	Male	Mean	SD	Mean	SD
Placebo	2	10	9.3	2.1	18.7	3.5
30	2	10	8.0	1.4	16.1	1.2
60	5	7	9.4	2.1	17.7	3.1
120	1	10	8.7	2.0	16.6	2.8
240	1	11	8.4	1.7	17.0	2.0
360	4	9	9.1	2.4	17.9	2.9
480	5	7	9.1	1.9	16.4	1.5

BMI, body mass index.

Table 2. Mean duration of urinary-concentrating action in hours stratified by gender.

			Placebo	30 µg	60 µg	120 µg	240 µg	360 µg	480 µg
DOA 125 mOsm/kg	n (censored ^a)	♀	2 (0)	2 (0)	5 (0)	1 (0)	1 (1)	4 (3)	5 (3)
		♂	10 (2)	10 (3)	7 (2)	10 (4)	11 (6)	9 (7)	7 (4)
	Mean (SD)	♀	0.0 (0.0)	2.0 (1.6)	4.9 (2.2)	7.6 ^b	11.6 ^b	9.4 (1.1)	11.0 (0.8)
		♂	0.7 (1.4)	3.9 (2.1)	6.5 (2.5)	8.2 (2.0)	9.5 (1.9)	10.6 (0.7)	10.2 (0.8)
	Gender difference	♂ - ♀	0.7	1.9	1.6	^c	^c	1.2	-0.8
DOA 200 mOsm/kg	n (censored ^a)	♀	2 (0)	2 (0)	5 (0)	1 (0)	1 (1)	4 (1)	5 (3)
		♂	10 (0)	10 (0)	7 (0)	10 (3)	11 (4)	9 (4)	7 (3)
	Mean (SD)	♀	0.0 (0.0)	0.9 (1.3)	3.8 (1.7)	7.2 ^b	9.8 ^b	8.8 (0.8)	9.8 (1.2)
		♂	0.1 (0.2)	2.7 (1.7)	6.1 (2.9)	7.5 (2.2)	8.9 (1.8)	9.5 (1.8)	9.7 (1.3)
	Gender difference	♂ - ♀	0.1	1.8	2.3	^c	^c	0.7	-0.1
DOA 400 mOsm/kg	n (censored ^a)	♀	2 (0)	2 (0)	5 (0)	1 (0)	1 (1)	4 (0)	5 (2)
		♂	10 (1)	10 (0)	7 (0)	10 (2)	11 (2)	9 (4)	7 (2)
	Mean (SD)	♀	0.0 (0.0)	0.0 (0.0)	2.9 (1.6)	3.5 ^b	9.8 ^b	8.0 (1.3)	8.9 (1.8)
		♂	0.0 (0.0)	1.6 (1.6)	4.7 (3.0)	6.1 (2.3)	8.1 (2.3)	8.6 (2.0)	8.5 (1.2)
	Gender difference	♂ - ♀	0	1.6	1.8	^c	^c	0.6	-0.4

^aPatients with osmolality above threshold level at the time their overhydration procedure stopped were censored; ^bSD not calculated as n = 1; ^cEstimates not calculated as n ≤ 1. OA, duration of urinary-concentrating action; SD, standard deviation.

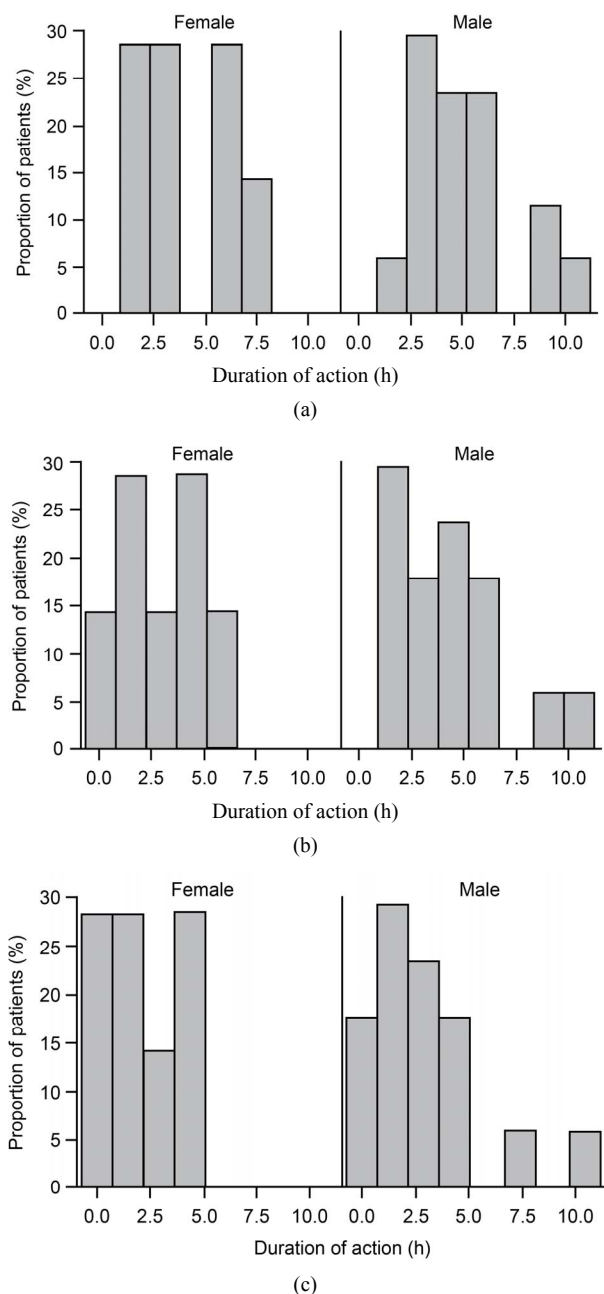


Figure 1. Distribution of duration of urine-concentrating action with urine osmolality above (a) 125 mOsm/kg, (b) 200 mOsm/kg and (c) 400 mOsm/kg in female and male patients following desmopressin 30 μ g or 60 μ g.

nary osmolality were similar in both boys and girls. Girls did not have a numerically higher mean, maximum or AUC of urinary osmolality following desmopressin administration compared with boys in any dose groups with sufficient sample size (**Table 3**). In the low-dose groups (desmopressin 30 and 60 μ g), boys tended to demonstrate a slightly greater PD response in terms of average, maximum and AUC of urinary osmolality compared with girls.

Plots of the distributions of time-adjusted AUC of osmolality, the AUC of urinary osmolality for ten hours following desmopressin administration and maximum urinary osmolality indicate that across all desmopressin doses tested boys had a slightly longer and more pronounced magnitude of antidiuretic response (**Figure 2**).

4. DISCUSSION

The small sample size and skewed gender distribution in this study prohibit a direct statistical comparison between genders. However, based on descriptive statistics, this *post hoc* analysis found no clear difference between the genders in either duration or magnitude of desmopressin antidiuretic action in most dose groups.

The most likely explanation for lack of a clear gender difference in children is that for most of the doses used in this study the upper asymptote of the dose-response curve for urine osmolality was already reached. In adults the desmopressin ODT ED_{50} for reduction in urine volume has been estimated to be 43.2 μ g and 16.1 μ g in men and women, respectively, using a maximal response (E_{max}) dose-response model with weight-adjusted dose (dose/weight) and a linear placebo effect [4]. The analysis estimated the dose sensitivity to desmopressin ODT for women relative to men is 2.7 (95% confidence interval 1.3 - 8.1) [4]. Taking into account the mean body weight of approximately 28 kg in this study, the likely exposure to desmopressin would be proportionally 2 - 3 times higher in children compared to adults [15]. Assuming linear extrapolation of desmopressin efficacy between adults and children, desmopressin ED_{50} values in children are likely to be approximately half the magnitude of ED_{50} values in adults. However, the desmopressin ED_{50} values for reduction in nocturnal urine volume estimated for adults were derived in patients with nocturia who had not been water-loaded and this study reports urine osmolality and DOA in water-loaded children following desmopressin administration. Nevertheless these data indicate that any future study exploring gender differences in the response to desmopressin ODT in children should include lower doses. We suggest that desmopressin ODT 10 μ g or a bioequivalent dose in other suitable formulations would be a suitable starting point.

Despite the ambiguity due to the skewed gender distribution, with more boys than girls, in the low-dose desmopressin ODT 30 and 60 μ g groups, boys had a 1 - 2 hours longer DOA compared with girls for all three threshold levels. A similar pattern was seen for mean, maximum and AUC of urinary osmolality, with boys receiving desmopressin ODT 30 and 60 μ g demonstrating higher values compared with girls. Although such a finding is contradictory to the expectation of greater female renal sensitivity to desmopressin, several studies

Table 3. Average, maximum and AUC of urinary osmolality stratified by gender.

			Placebo	30 µg	60 µg	120 µg	240 µg	360 µg	480 µg
n		♀	2	2	5	1	1	4	5
		♂	10	10	7	10	11	9	7
Average osmolality during action (mOsm/kg)	Mean (SD)	♀	37.2	136.5	381.2	569.3	844	670.2	621.9
		♂	51.9	305.6	524.4	576.8	656.8	684.1	665.5
	Gender difference	♂ - ♀	14.7	169.1	143.2	^a	^a	13.9	43.6
Maximum osmolality during action (mOsm/kg)	Mean (SD)	♀	83.5	263	777.2	935	1049	976.8	891.8
		♂	179.7	565.2	823.7	894.9	948.37	931.1	978.7
	Gender difference	♂ - ♀	96.2	302.2	46.5	^a	^a	-45.7	86.9
AUC of osmolality 0 - 10 hours (mOsm/kg)	Mean (SD)	♀	278.1	775.5	3047.3	4844.7	8301	7079.3	6425.5
		♂	430.5	1932.6	4152	5352.2	6609.3	7085.2	6455.8
	Gender difference	♂ - ♀	152.4	1157.1	1104.7	^a	^a	5.9	30.3

^aEstimates not calculated as $n \leq 1$. AUC, area under the curve; SD, standard deviation.

have demonstrated greater male sensitivity to AVP in young rats [16-18]. One study comparing the renal sensitivity to AVP and density of V2 receptors in young female and male rats (aged 2 - 3 months) found that the young male rats had significantly greater sensitivity to AVP infusion and also significantly greater density of V2 receptors than young female rats [19]. However, a similar study in adult rats found the opposite result—female rats expressed significantly more V2 receptor messenger RNA and protein in the kidney than male rats, resulting in a greater sensitivity to V2 receptor agonist administration [20]. Thus, as there are no obvious reasons for the contradictory findings in these rat studies other than the different ages and life-stages of the animals, it suggests that sensitivity to V2 receptor stimulation is greater in juvenile male individuals, and this inverts to greater female sensitivity with sexual maturity and the influence of oestrogen, known to influence osmoregulation [21], and other sex hormones. Some pre-clinical studies have suggested that escape from X inactivation may alter with developmental stages or ageing, providing a possible mechanism for increased female V2R expression with age [22,23].

A gender difference in prevalence of MNE has been shown, with twice as many boys as girls affected by the condition [7]. Furthermore, the normal circadian increase in nocturnal AVP levels is absent in younger patients with enuresis [24]. However, it has not been reported whether puberty and increased levels of endogenous sex hormones increase AVP sensitivity to a larger degree in girls compared with boys. If our finding of no gender difference in response to V2 receptor stimulation in the paediatric population can be confirmed in a prospective study using lower doses than in our study, this will support the hypothesis that we proposed previously [4]; a gender difference in sensitivity to desmopressin is absent

in children since sex hormones are at low levels prior to puberty.

Another possibility for lack of gender difference in children is the recently discovered age-dependent skewing of X chromosome inactivation [25]. While in general young females have a mosaic of cells with either the maternal or the paternal X chromosome inactivated with a 50:50 distribution, a deviation from this balanced expression may occur with ageing [25]. This may clinically manifest as late-onset X-linked disorders. The AVPR2 gene has known gain and loss of function mutations, leading to syndrome of inappropriate antidiuresis and nephrogenic diabetes insipidus (NDI), respectively [26]. A case of transient NDI associated with decreased V2 receptor expression has been reported in a 55-year-old woman following surgery [27]. However, it remains to be shown if the AVPR2 gene located at xq-28 is subject to late-onset skewing of inactivation, how common such skewing might be and whether other mutations may contribute to V2 receptor expression differences.

Comparative pharmacokinetic (PK)/PD studies of desmopressin have only been performed in a paediatric population by Fjellestad-Paulsen *et al.* and no gender differences were reported [28]. As in our study, the study populations were predominantly male, and sample sizes were too small to allow gender stratification.

Further research is therefore needed to explore potential gender differences in response to desmopressin in a prospective setting, including sufficient paediatric patients of both genders to allow for a stratified analysis. The current analysis supports power calculations for any such prospective study protocols in a paediatric population on gender difference in response to V2 receptor stimulation. Furthermore, to reduce variation in PD endpoints due to PK variation, we recommend such a study to be conducted by intravenous infusion of desmopressin.

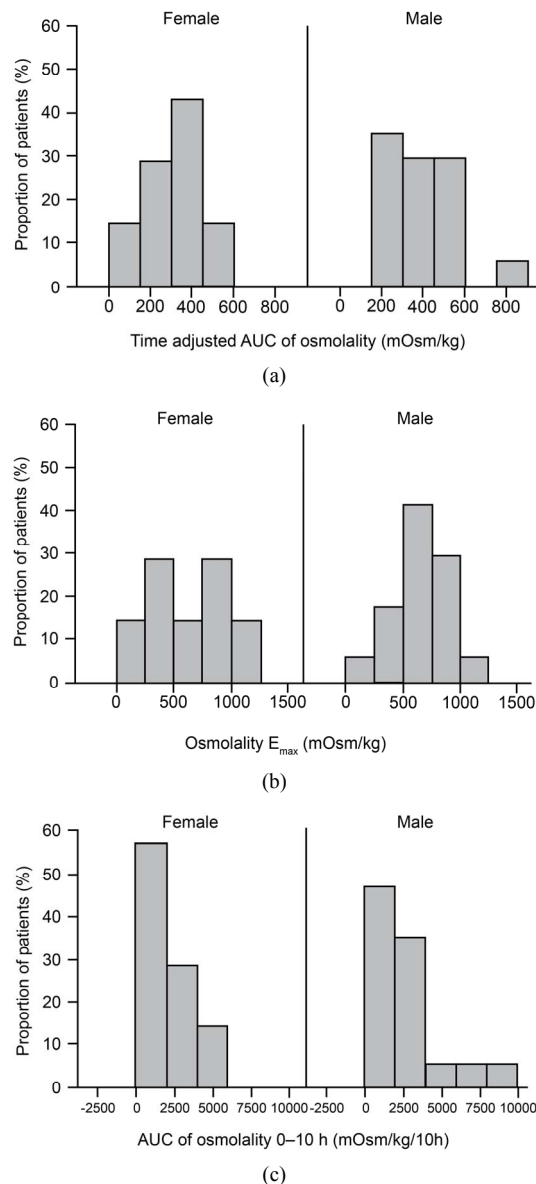


Figure 2. Distribution of urine osmolality reported as time adjusted AUC, (b) E_{max} and (c) AUC over 0 - 10 hours in female and male patients following desmopressin 30 - 480 µg. AUC, area under the curve.

A previous study designed to investigate gender differences in desmopressin response in young adults used a bolus injection [29], which does not mimic steady state conditions, but only briefly saturates the V2 receptors. Therefore, we suggest a more suitable approach is to infuse continuously a low dose of desmopressin intravenously (to reach similar concentrations to those achieved using desmopressin ODT 10 µg) until steady state is reached and then gradually increase the dose.

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6. CONFLICT OF INTEREST/DISCLOSURE

Kristian Vinter Juul, Sandra Goble and Jens Peter Nørgaard are employees of Ferring Pharmaceuticals.

Johan Vande Walle has acted as a safety board adviser for Ferring International.

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